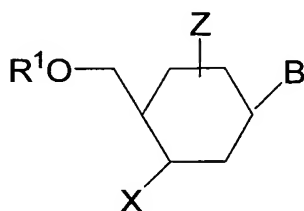


AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims

1. (Currently Amended): A six membered, at least partially unsaturated, carbocyclic nucleoside compound, including the (-) enantiomer, the (+) enantiomer, and pharmaceutically acceptable salts and esters thereof, the compounds represented by formula I:



I

wherein:

-Z represents the presence of one double bond in the six membered carbocyclic ring,

-B is a heterocyclic ring selected from the group consisting of pyrimidine and purine bases,

-X is an azido, F, or OR²,

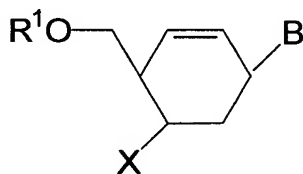
-R¹ and R² are the same or different and represent the same or different protecting groups which are combined to form a protecting group or are each a protective group, hydrogen, alkyl, alkenyl, acyl or phosphate moieties wherein;

-the alkyl moiety is a saturated, substituted or unsubstituted straight or branched ~~chains~~ chain hydrocarbon radical having from 1 to 20, ~~1-16, 1-14, 1-12, 1-10, 1-8,~~
~~1-4,~~ carbon atoms,

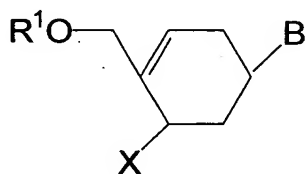
-the alkenyl moiety is an unsaturated congener of the alkyl group and,

-the acyl moiety is an alkanoyl or aroyl moiety, wherein alkanoyl is an alkyl carbonyl radical, wherein alkyl is as described above and aroyl represents benzoyl substituted benzoyl or naphthoyl.

2. (Previously Presented): A six membered, at least partially unsaturated, carbocyclic nucleoside compound, according to claim 1, being a cyclohexenyl nucleoside compound having a formula selected from the group consisting of II and III:

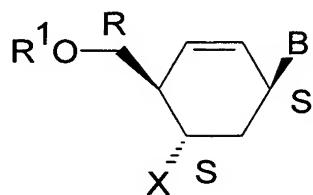


II

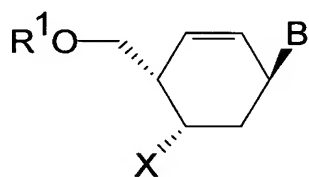


III

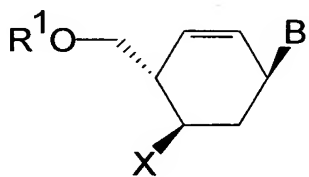
3. (Previously Presented): Compound according to claim 1, selected from the group of compounds consisting of IV, V, VI, VII, VIII, IX, X and X':



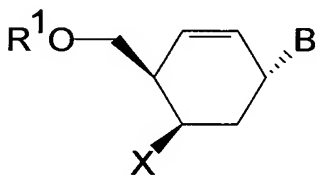
IV



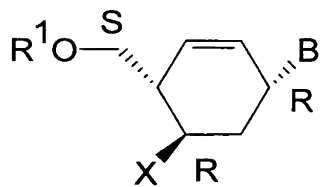
V



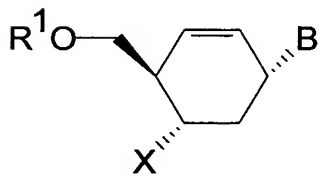
VI



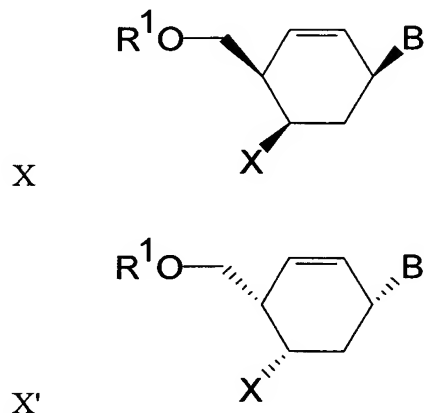
VII



VIII



IX



4. (Previously Presented): Compound according to claim 1, wherein the C₁ bearing B substituent and the C₅ bearing X substituent both have the (S)-configuration, and the C₄ bearing –OR¹ substituent has the (R)-configuration, as depicted by formula IV in claim 3.

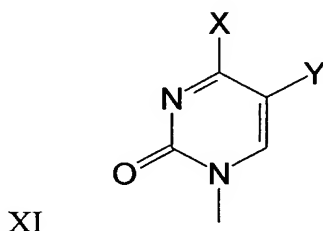
5. (Previously Presented): Compound according to claim 1, wherein the C₁ bearing B substituent and the C₅ bearing X substituent both have the (R)-configuration, and the C₄ bearing –OR¹ substituent has the (S)-configuration, as depicted by formula VIII in claim 3.

6. (Previously Presented) Compound according to claim 1, wherein X is represented by a hydroxyl group in the (S)-configuration.

7. (Previously Presented) Compound according to claim 1, wherein X is hydroxyl in the (R)-configuration.

8. (Withdrawn) Compound according to claim 1, wherein B is derived from the group consisting of pyrimidine bases.

9. (Withdrawn): Compound according to claim 7, wherein the pyrimidine base has formula XI:



wherein X is chosen from the group consisting of:

- OH, NH₂, and NHQ,

wherein;

- Q is selected from the group consisting of:

OH and C₁₋₅ alkyl, and

- Y is selected from the group consisting of:

H, F, Cl, Br, I, C₁₋₅ alkyl, haloethyl and CH=CH-R, wherein R represents hydrogen, halogen or C₁₋₅ alkyl, and wherein haloethyl contains from 1 to 4 F, Cl or Br atoms.

10. (Withdrawn): Compound according to claim 1, wherein B is selected from the group consisting of purine bases which are optionally substituted with aza, deaza, deoxy or deamino analogues, guanine, 2,6-diaminopurine, hypoxanthine and xanthine.

11. (Withdrawn): Compound according to claim 1, wherein the B is selected from the group consisting of aza, deaza, deoxy and deamino analogues of the heterocyclic rings.

12. (Previously Presented): Compound according to claim 1, wherein the protecting group is selected from the group consisting of a silyl protecting group, a benzyl protecting group, a benzoyl protecting group and a $C_6H_5-CH=$ group.

13. (Withdrawn): Compound according to claim 1, selected from the group consisting of:

- 9-[1S,4R,5S)-5-hydroxy-4-hydroxymethyl-2-cyclohexenyl] guanine, and
- 9-[1R,4S,5R)-5-hydroxy-4-hydroxymethyl-2-cyclohexenyl].

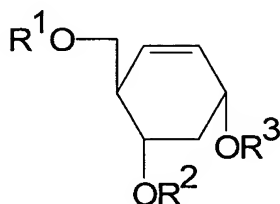
14. (Previously Presented) Compound according to claim 1 selected from the group consisting of:

- 9-[(1S,4R,5S)-5-(tert-Butyldimethylsilyloxy)-4-(tert-butyldimethylsilyloxymethyl)-2-cyclohexenyl] adenine
- 9-[(1S,4R,5S)-5-Hydroxy-4-hydroxymethyl-2-cyclohexenyl]adenine
- 9-[(1S,4R,5S)-5-(tert-butyldimethylsilyloxy)-4-(tert-butyldimethylsilyloxymethyl)-2-cyclohexenyl]-2-amino-6-chloropurine
- 9-[(1S,4R,5S)-5-hydroxy-4-hydroxymethyl-2-cyclohexenyl]guanine
- 9-[(1R,4S,5R)-5-Benzoyloxy-4-benzoyloxymethyl-2-cyclohexenyl] adenine
- 9-[(1R,4S,5R)-5-hydroxy-4-hydroxymethyl-2-cyclohexenyl] adenine

- 9-[(1R,4S,5R)-5-Benzoyloxy-4-benzoyloxymethyl-2-cyclohexenyl]
guanine, and
- 9-[(1R,4S,5R)-5-Hydroxy-4-hydroxymethyl-2-cyclohexenyl] guanine.

15. (Currently Amended): A method of producing the compound of claim 1, including, the (-) enantiomer, the (+) enantiomer, and pharmaceutically acceptable salts and esters thereof, comprising the steps of:

- i) ~~reacting~~ providing a cyclohexenyl compound of formula XII:



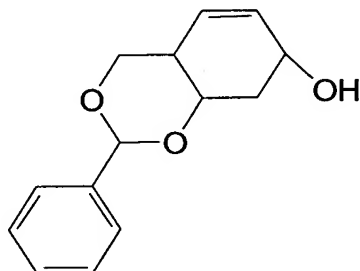
- ii) substituting the OR³-group with a purine base.

wherein R¹ and R² is are combined to form a protecting group or are each protecting groups and R³ is a leaving group or a Hydrogen atom, and wherein the OR³-group of the cyclohexenyl compound is substituted by a pyrimidine or purine base.

16. (Previously Presented): Process according to claim 15 wherein R³ is hydrogen and wherein a Mitsunobo reaction is utilised.

17. (Previously Presented): Process according to claim 15 wherein R³ is a leaving group enabling nucleophilic substitution.

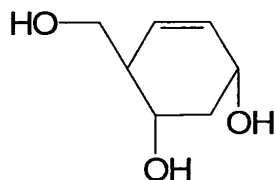
18. (Previously Presented): Process according to claim 15, wherein the compound of formula XII has the chemical formula XIII;



XIII

including analogues thereof either in a racemate form or separated isomers thereof.

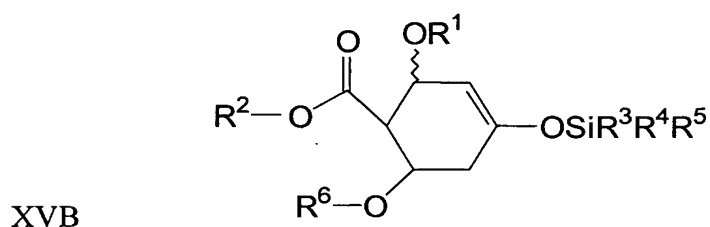
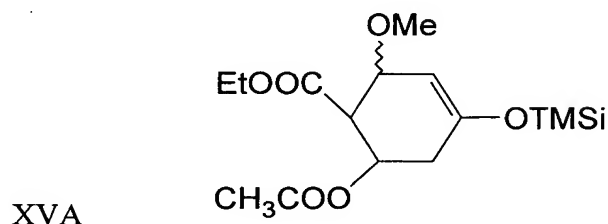
19. (Previously Presented): Process according to claim 18, wherein compound XIII is provided by reacting (\pm) 4-hydroxymethyl-cyclohex-2-en-1,5 Diol of formula XIV;



XIV

with a benzaldehyde analogue and a Lewis acid.

20. (Previously Presented): Process according to claim 15, wherein compound XIV is provided by the reduction of compound selected from the group consisting of XVA and XVB;



wherein for XVB:

- R^1 and R^2 are alkyl or alkenyl moieties,

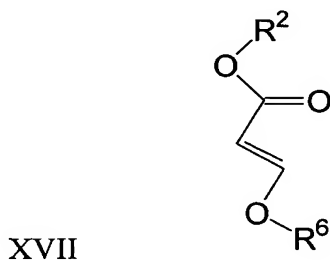
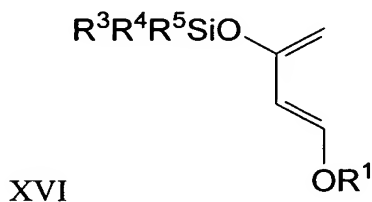
wherein:

- R^1 and R^2 are the same or different, and
- alkyl is a saturated, substituted or unsubstituted hydrocarbon radical having from 1 to 20, carbon atoms and being straight or branched chain, and
- alkenyl is the unsaturated congener of the alkyl group, and
- R^3 , R^4 and R^5 are alkyl, alkenyl or aryl moieties, wherein:
 - R^3 , R^4 and R^5 are the same or different, and
 - alkyl is a saturated, substituted or unsubstituted straight or branched chain hydrocarbon radical having from 1 to 20 carbon atoms and
 - alkenyl is the unsaturated congener of the alkyl group, and
 - aryl represents phenyl or substituted phenyl, and
- R^6 is an alkyl, alkenyl or acyl moiety, wherein

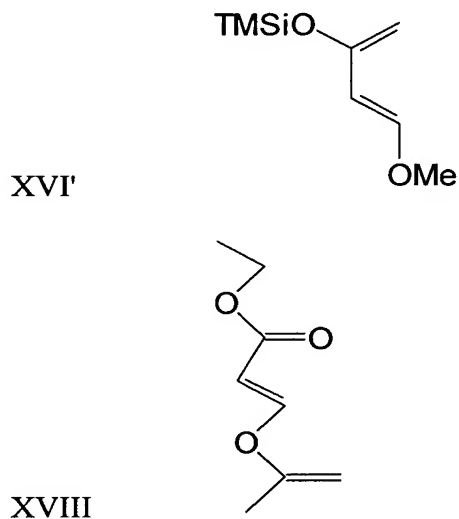
- alkyl is a saturated, substituted or unsubstituted hydrocarbon straight or branched chain radical having from 1 to 20 carbon atoms,
- alkenyl is the unsaturated congener of the alkyl group, and
- acyl is an alkanoyl or aroyl moiety, wherein alkanoyl is an alkyl carbonyl radical, wherein alkyl is as described above and aroyl represents benzoyl, substituted benzoyl or naphthoyl.

21. (Previously Presented): Process according to claim 20, wherein compound XVA or XVB is provided by a Diels-Alder reaction, by the cyclo addition of a suitable diene and dienophile.

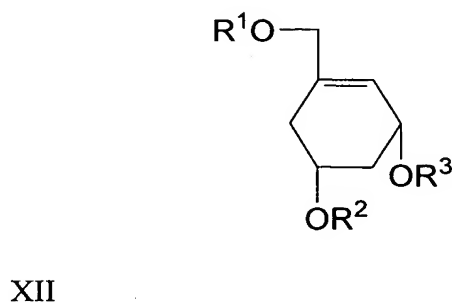
22. (Previously Presented): Process according to claim 21 wherein the diene has the following chemical structure XVI, and the dienophile has the following chemical structure XVII, wherein R^1 , R^2 , R^3 , R^4 , R^5 and R^6 are as defined in claim 20;

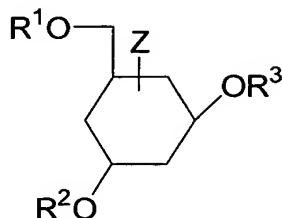


23. (Previously Presented): Process according to claim 22 wherein the diene has the chemical structure XVI' and the dieneophile has the chemical structure XVIII;



24. (Withdrawn): A six membered, at least partially unsaturated, carbocyclic nucleoside compound, including the (-) enantiomer, the (+) enantiomer, and pharmaceutically acceptable salts and esters thereof, the compounds represented by a formula selected from the group consisting of XII and XIX;





XIX

wherein:

- Z represents the presence of one double bond in the carbocyclic ring,
- R¹ and R² are protecting groups and R³ is a leaving group or a Hydrogen atom.

25. (Withdrawn): Compound according to claim 24, wherein:

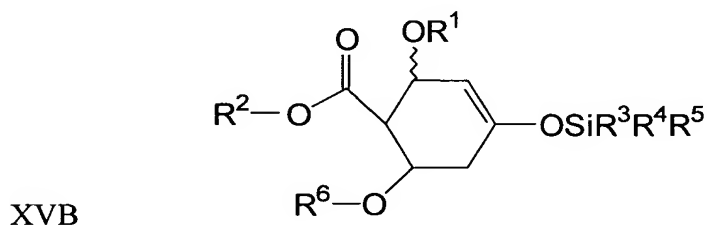
R¹ and R² are the same or different and hydrogen, alkyl, alkenyl, acyl or phosphate moieties are represented, or R¹ and R² represent a cyclic protecting group, wherein:

- alkyl is a saturated, substituted or unsubstituted straight or branched chain hydrocarbon radical having from 1 to 20 carbon atoms, and
- alkenyl is the unsaturated congener of the alkyl group, and
- acyl is an alkanoyl or aroyl moiety, wherein alkanoyl is an alkyl carbonyl radical, wherein alkyl is as described above and aroyl represents benzoyl, substituted benzoyl or naphthoyl; and

R^3 represents a hydrogen, an alkylsulfonyl or an arylsulfonyl moiety, wherein:

- alkyl is a saturated, substituted or unsubstituted hydrocarbon radical having from 1 to 6 carbon atoms and straight or branched chain, and
- aryl represents phenyl or substituted phenyl.

26. (Withdrawn): A cyclohexenyl compound, including the (-) enantiomer, the (+) enantiomer, and pharmaceutically acceptable salts and esters thereof, the compound represented by formula XVB;



wherein R^1 and R^2 are alkyl or alkenyl moieties, wherein

- R^1 and R^2 are the same or different, and
- alkyl is a saturated, substituted or unsubstituted straight or branched chain hydrocarbon radical having from 1 to 20 carbon atoms,
- alkenyl is the unsaturated congener of the alkyl group, and

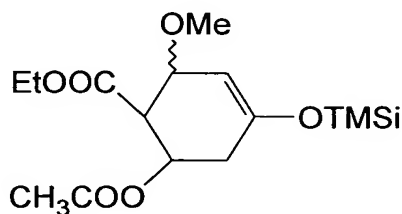
R^3 , R^4 and R^5 are alkyl, alkenyl or aryl moieties, wherein:

- R^3 , R^4 and R^5 are the same or different, and
- alkyl is a saturated, substituted or unsubstituted straight or branched chain hydrocarbon radical having from 1 to 20 carbon atoms and,
- alkenyl is the unsaturated congener of the alkyl group, and
- aryl represents phenyl or substituted phenyl, and

R^6 is an alkyl, alkenyl or acyl moiety, wherein:

- alkyl is a saturated, substituted or unsubstituted straight or branched chain hydrocarbon radical having from 1 to 20 carbon atoms, and
- alkenyl is the unsaturated congener of the alkyl group, and
- acyl is an alkanoyl or aroyl moiety, wherein alkanoyl is an alkyl carbonyl radical, wherein alkyl is as described above and aroyl represents benzoyl, substituted benzoyl or naphthoyl.

27. (Withdrawn): Compound according to claim 26 having the formula XVA being; 5-O-acetyl-4-ethoxycarbonyl-3-O-methyl-1-O-trimethylsilyl-cyclohexen-1,3,5-triol and its isomers;



XVA

28. (Withdrawn): A compound having the formula (\pm) 4-hydroxymethyl-cyclohex-2-en-1,5-diol.

29. (Withdrawn): A compound having the formula (1R, 4R, 5S)-4-hydroxymethyl-cyclohex-2-en-1,5-diol.

30. (Withdrawn): A compound having the formula (1S, 4S, 5R)-4-hydroxymethyl-cyclohex-2-en-1,5-diol.

31. (Withdrawn): A compound having the formula (\pm) 5,7-O-benzylidene-4-hydroxymethyl-cyclohex-2-en-1,5-diol.

32. (Withdrawn): A compound having the formula (1R, 4R, 5S)-5,7-O-benzylidene-4-hydroxymethyl-cyclohex-2-en-1,5-diol.

33. (Withdrawn): A compound having the formula (1S, 4S, 5R) 5,7-O-benzylidene-4-hydroxymethyl-cyclohex-2-en-1,5-diol.

34. (Withdrawn): Compound according to claim 24 selected from the group consisting of:

- (4S,5R)-5-Benzoyloxy-4-benzoyloxymethyl-cyclohex-2-en-1-one,
- (1S,4S,5R)-5-Benzoyloxy-4-benzoyloxymethyl-cyclohex-2-en-1-ol,
- (4R,5S)-4-tert-Butyldimethylsilyloxymethyl-5-tert-butyldimethylsilyloxy-cyclohex-2-en-1-one, and
- (1R,4R,5S)-5-(tert-Butyldimethylsilyloxy)-4-(tert-butyldimethylsilyloxymethyl)-cyclohex-2-en-1-ol.

35. (Previously Presented): Compound obtained by the process of claim 15.

36. ((Previously Presented): Pharmaceutical composition comprising a compound and a carrier according to claim 1.

37. (Previously Presented): A pharmaceutical composition as claimed in claim 1, having antiviral activity towards herpetic viruses.

38. (Previously Presented): A pharmaceutical composition as claimed in claim 37 comprising said active ingredient in a concentration ranging from about 0.1-100 % by weight.

39. (Previously Presented): A pharmaceutical composition as claimed in claim 38, having a form which is selected from the group consisting of powders, suspensions, solutions, sprays, emulsions, unguents and creams.

40. (Canceled).

41. (Previously Presented): A method of providing antiviral biological activity against herpes viruses, pox viruses and related viruses, comprising administering the compound according to claim 1.

42. (Canceled).

43. (Previously Presented): A method for the preparation of a pharmaceutical composition having antiviral activity against herpes viruses, pox viruses and related viruses, comprising combining the compound according to claim 1 with other ingredients.

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Response to Office Action dated September 26, 2005
Paper dated October 23, 2006
Attorney Docket No. 0702-020249

44. (Previously Presented): Method of claim 41, wherein the biological activity comprises pharmaceutical activity.

45. (Canceled)